## Amendments to the Claims

Following is a complete set of claims as amended with this Response. This complete set of claims excludes cancelled claim 25 and includes amended claims 26, 32, and 34.

(Withdrawn) A method of performing electrophysiological testing in a 1. cardiac stimulation device capable of delivering non-invasive programmed stimulation, comprising:

detecting a cardiac event in a cardiac chamber;

implementing an electrophysiological testing scheme upon detection of the cardiac event occurring in the cardiac chamber; and

delivering a predetermined sequence of stimulation pulses to the cardiac chamber as dictated by the testing scheme.

- (Withdrawn) The method of claim 1, wherein implementing the testing 2. scheme is performed during a refractory period that follows the detected cardiac event.
- (Withdrawn) The method of claim 2, wherein implementing the testing 3. scheme includes switching from a standard operating mode to a non-invasive programmed stimulation mode.
- (Withdrawn) The method of claim 3, further including receiving an 4. external command that triggers the onset of the non-invasive programmed stimulation.
- (Withdrawn) The method of claim 3, wherein detecting the cardiac event 5. includes detecting an intrinsic event in the cardiac chamber being tested.
- (Withdrawn) The method of claim 5, wherein detecting an intrinsic event 6. includes detecting an intrinsic depolarization occurring in one of an atrial cardiac chamber and a ventricular cardiac chamber.

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- 7. (Withdrawn) The method of claim 3, wherein detecting the cardiac event includes detecting a stimulated event in the cardiac chamber being tested.
- 8. (Withdrawn) The method of claim 7, wherein detecting a stimulated event includes detecting one of an atrial stimulation pulse and a ventricular stimulation pulse.
- 9. (Withdrawn) The method of claim 3, further including providing a recovery delay following the non-invasive programmed stimulation.
- 10. (Withdrawn) The method of claim 9, further comprising starting a second refractory period following the expiration of the recovery delay if no intrinsic event is detected during the recovery delay.
- 11. (Withdrawn) The method of claim 10, further including effecting a transfer from the non-invasive programmed stimulation mode to the standard operating mode during the second refractory period.
- 12. (Withdrawn) The method of claim 1, further including blanking sensing circuitry of non-tested cardiac chambers during the delivery of the sequence of stimulation pulses in the cardiac chamber being tested.
- 13. (Withdrawn) The method of claim 1, further including providing back-up ventricular stimulation whenever atrial non-invasive programmed stimulation is performed; and

wherein providing back-up ventricular stimulation includes providing backup ventricular stimulation at a programmed rate that is decoupled from the atrial non-invasive programmed stimulation.

- (Withdrawn) The method of claim 9, further comprising starting a 14. refractory period if an intrinsic event is sensed in the recovery period.
- (Original) A stimulation device capable of performing electrophysiological 15. testing by delivering non-invasive programmed stimulation, comprising:

a discriminator that senses a cardiac event in a cardiac chamber being tested:

timing circuitry, coupled to the discriminator, that triggers an onset of the non-invasive programmed stimulation based on a detected cardiac event occurring in the cardiac chamber being tested;

a controller, connected to the timing circuitry that executes a transfer between a first and a second stimulation mode; and

an energy generator connected to the discriminator, the timing circuitry and the controller, the generator is controlled by the controller to deliver a sequence of stimulation pulses to the cardiac chamber being tested in response to the detected cardiac event.

- (Original) The stimulation device of claim 15, wherein the timing circuitry 16. sets a refractory period that follows a triggering detected cardiac event; and wherein the controller executes the transfer during the refractory period.
- (Original) The stimulation device of claim 16, wherein the controller 17. executes the transfer between the first and the second stimulation mode by switching from a standard operating mode to a non-invasive programmed stimulation mode.
- (Original) The stimulation device of claim 17, further including a 18. programmer that generates an external command; and

wherein the timing circuitry triggers the onset of the non-invasive programmed stimulation in response to the external command.

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- (Original) The stimulation device of claim 17, wherein the discriminator 19. detects any one of an atrial intrinsic event, ventricular intrinsic event, an atrial stimulated event, or a ventricular stimulated event in the cardiac chamber being tested.
- (Original) The stimulation device of claim 17, wherein the timing circultry 20. further sets a recovery delay at the expiration of the non-invasive programmed stimulation.
- (Original) The stimulation device of claim 20, wherein the timing circuitry 21. is operative to start a second refractory period following the expiration of the recovery delay if no intrinsic event is detected during the recovery delay.
- (Original) The stimulation device of claim 21, wherein the controller 22. further effects a transfer from the non-invasive programmed stimulation mode to the standard operating mode during the second refractory period.
- (Original) The stimulation device of claim 15, wherein the energy 23. generator further provides back-up ventricular stimulation whenever atrial non-invasive programmed stimulation is performed.
- (Original) The stimulation device of claim 23, wherein the energy 24. generator provides back-up ventricular stimulation at a programmed rate that is decoupled from the atrial non-invasive programmed stimulation.
  - (Currently Cancelled) 25.

- 26. (Currently Amended) The stimulation device of claim 25, A stimulation device capable of performing electrophysiological testing by delivering non-invasive programmed stimulation, comprising:
  - a sensing circuitry to detect a cardiac event in a cardiac chamber to be tested;
  - a controller coupled to the sensing circuitry, the controller to implement an electrophysiological testing scheme in response to detection of the cardiac event; and
- a pulse generator coupled to the controller, the pulse generator to deliver a sequence of stimulation pulses to the cardiac chamber as dictated by the testing scheme;

wherein the controller comprises a timing control circuitry coupled to the sensing circuitry, the timing control circuitry to trigger an onset of the non-invasive programmed stimulation based on the detected cardiac event occurring in the cardiac chamber being tested; and

wherein the controller implements the testing scheme during a refractory period.

- 27. (Previously Presented) The stimulation device of claim 26, wherein the electrophysiological testing scheme comprises a transfer from a standard operating mode to a non-invasive programmed stimulation mode.
- 28. (Previously Presented) The stimulation device of claim 27, wherein the sensing circuitry detects any one of an atrial intrinsic event, ventricular intrinsic event, an atrial stimulated event, or a ventricular stimulated event in the cardiac chamber being tested.
- 29. (Previously Presented) The stimulation device of claim 27, wherein the timing control circuitry further sets a recovery delay at the expiration of the non-invasive programmed stimulation.

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- 30. (Previously Presented) The stimulation device of claim 29, wherein the timing control circuitry is operative to start a second refractory period following the expiration of the recovery delay if no intrinsic event is sensed during the recovery delay.
- 31. (Previously Presented) The stimulation device of claim 30, wherein the controller further effects a transfer from the non-invasive programmed stimulation mode to the standard operating mode during the second refractory period.
- 32. (Currently Amended) The stimulation device of claim 25 26, wherein the pulse generator further provides back-up ventricular stimulation whenever atrial non-invasive programmed stimulation is performed.
- 33. (Previously Presented) The stimulation device of claim 32, wherein the pulse generator provides back-up ventricular stimulation at a programmed rate that is decoupled from the atrial non-invasive programmed stimulation.
- 34. (Currently Amended) The stimulation device of claim 25 26, wherein the controller further effects a transfer from the test mode to a normal mode if a failure occurs during the non-invasive programmed stimulation.
  - 35. (Previously Cancelled)